In the Claims

Please amend the claims as follows:

Claim 1 (Previously Amended) A compound of Formula I

or a pharmaceutically acceptable salt of said compound; wherein R_1 is a) -(C_1 - C_6)alkyl optionally substituted with -CF₃, b) -C=C-CH₃, c) -C=C-CI, d) -C=C-CF₃, e) -CH₂O(C_1 - C_4)alkyl optionally substituted with -CF₃ or f) -CF₃;

 $R_2 \text{ is a) -(C_1-C_5)alkyl, b) -(C_2-C_5)alkenyl or c) -phenyl optionally substituted} \\ \text{with one of the following: -OH, -NR_9-C(O)-(C_2-C_4)alkyl, -CN, -Z-het, -O-(C_1-C_3)alkyl-C(O)-NR_9R_{10}, -NR_9-Z-C(O)-NR_9R_{10}, -Z-NR_9-SO_2-R_{10}, -NR_9-SO_2-het, -O-C(O)-(C_1-C_4)alkyl or -O-SO_2-(C_1-C_4)alkyl;} \\$

Z for each occurrence is independently $-(C_0-C_4)$ alkyl;

 R_3 is a) -hydrogen, b) -(C_1 - C_6)alkyl optionally substituted with one to three halo, c) -(C_2 - C_6)alkenyl or d) -(C_2 - C_6)alkynyl optionally substituted with one to three halo;

 R_4 is a) -hydrogen, or b) -(C_2 - C_5)alkyl-NR₅R₆;

R₅ and R₆ are each independently a) hydrogen or b) -(C₁-C₃)alkyl;

het is an optionally substituted 5-, 6- or 7-membered saturated, partially saturated or unsaturated heterocyclic ring containing from 1 to 3 heteroatoms selected from the group consisting of nitrogen, oxygen and sulfur; and including any bicyclic group in which any of the above heterocyclic rings is fused to a benzene ring or another heterocyclic ring; and optionally

substituted with one to four R₇; provided that het is other than pyridinyl, imidazolyl or tetrazolyl;

 R_7 is a) -(C_1 - C_6)alkyl optionally substituted with one to three R_8 , b) –Z- NR_9R_{10} or c) –Z-C(O)- NR_9R_{10} ;

 R_8 for each occurrence is independently a) halo, b) –OH, c) oxo or d) - O(C_1 - C_6)alkyl;

 R_9 and R_{10} for each occurrence are independently a) -H or b) -(C_1 - C_3)alkyl; or R_9 and R_{10} are taken together with N to form het; provided that:

- 1) when R_1 is $-C \equiv C CH_3$, R_2 is phenyl and R_3 is hydrogen, then R_4 is other than $-(CH_2)_2 N(CH_3)_2$, or $-(CH_2)_3 N(CH_3)_2$, $-(CH_2)_2$ -pyrrolidinyl optionally substituted with methyl, $-(CH_2)_3$ -pyrrolidinyl or $-(CH_2)_2$ -morpholinyl;
- 2) when R_1 is $-C \equiv C CH_3$, R_2 is propyl and R_3 is hydrogen, then R_4 is other than $-(CH_2)_2 N(CH_3)_2$; and
- 3) when R_1 is $-C \equiv C CH_3$, R_2 is butyl and R_3 is hydrogen, then R_4 is other than $-(CH_2)_2 N(CH_3)_2$.

Claim 2 (Previously Amended) A compound of claim 1 of Formula II

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or a pharmaceutically acceptable salt of said compound;

wherein R_1 is a) -(C_1 - C_6)alkyl optionally substituted with - CF_3 , b) - $C \equiv C$ - CH_3 , c) - CF_3 or d) -- $CH_2O(C_2$ - C_4)alkyl.

Claim 3 (Original) A compound of claim 2 wherein R_1 is a) $-CH_2CH_2CH_3$, b) $-C = C - CH_3$ or c) $-CF_3$.

Claim 4 (Original) A compound of claim 3 wherein R₃ is a) hydrogen, b) methyl, c) ethyl, d) propyl or e) isopropyl;

 R_4 is -(C_2 - C_3)alkyl-NR₅R₆;

 R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim 5 (Original) A compound of claim 4

wherein R₃ is a) methyl, b) ethyl, c) propyl or d) isopropyl;

 R_4 is -(C_2 - C_3)alkyl-NR₅R₆;

 R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim 6 (Original) A compound of claim 5

wherein R₃ is a) methyl or b) ethyl;

 R_4 is -(C_2 - C_3)alkyl-NR₅R₆;

 R_5 and R_6 are each methyl.

Claims 7-11 (Previously Canceled)

Claim 12 (Original) A compound of claim 1

wherein R_1 is a) $-CH_2CH_2CH_3$, b) $-C = C-CH_3$ or c) $-CF_3$;

 R_2 is a) -(C_1 - C_5)alkyl or b) -(C_2 - C_5)alkenyl;

R₃ is a) hydrogen, b) methyl, c) ethyl, d) propyl or e) isopropyl;

 R_4 is -(C_2 - C_3)alkyl-NR₅R₆;

 R_5 and R_6 are each independently a) methyl, b) ethyl, c) propyl or d) isopropyl.

Claim13 (Original) A compound of claim 12 wherein R₂ is a) methyl, b) ethyl, c) propyl, d) ethenyl, e) propenyl or f) butenyl;

R₃ is a) hydrogen, b) methyl or c) ethyl,

R₅ and R₆ are each independently a) methyl or b) ethyl.

Claims 14-17 (Previously Canceled)

Claim 18 (Original) A compound of claim 1 wherein in Formula I –CH₂-R₂ is ethenyl or ethynyl.

Claim 19 (Original) A compound of claim 4 selected from the group consisting of:

carbamic acid, [2-(dimethylamino)ethyl]-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester;

carbamic acid, [3-(dimethylamino)propyl]-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester; and

carbamic acid, [3-(diethylamino)propyl]-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester.

Claim 20 (Original) A compound of claim 6 selected from the group consisting of:

carbamic acid, [2-(dimethylamino)ethyl]methyl-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester;

carbamic acid, [2-(dimethylamino)ethyl]methyl-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-propyl-2-phenanthrenyl ester;

carbamic acid, [3-(dimethylamino)propyl]ethyl-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester; and

carbamic acid, [2-(dimethylamino)ethyl]ethyl-, (4bS,7R,8aR)-4b,5,6,7,8,8a,9,10-octahydro-7-hydroxy-4b-(phenylmethyl)-7-(trifluoromethyl)-2-phenanthrenyl ester.

Claim 21-23 (Previously Canceled)

Claim 24 (Original) A compound of claim 13 selected from the group consisting of:

carbamic acid, (3-dimethylaminopropyl)methyl-, (4bS, 7R, 8aR)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester;

carbamic acid, (2-dimethylaminoethyl)methyl-, (4bS, 7R, 8aR)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester;

carbamic acid, (2-dimethylaminoethyl)ethyl-, (4bS, 7R, 8aR)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester; and

carbamic acid, (2-dimethylaminoethyl)-, (4bS, 7R, 8aR)-4b,5,6,7,8,8a,9,10-octahydro-4b-ethyl-7-hydroxy-7-prop-1-ynyl-phenanthren-2-yl ester.

Claims 25-26 (Previously canceled)

Claim 27 (Newly amended) A method for the treatment of a glucocorticoid receptor-mediated disease or condition which is selected from the group consisting of obesity, diabetes, depression, anxiety and neurodegeneration in a mammal, which comprises administering to the mammal a therapeutically effective amount of a compound of claim 1, or a pharmaceutically acceptable salt of said compound.

Claim 28 (Previously canceled)

Claim 29 (Previously amended) The method of claim 27 wherein the condition is obesity.

Claims 30-41 (Previously canceled)